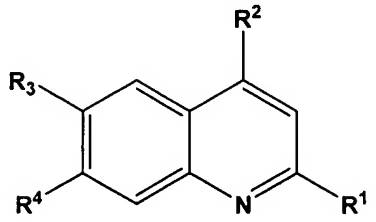
Amendments to the Claims:

This listing of claims will replace all prior versions and listing of claims in the application.

Please amend claims 1, 3 to 6, 9 and 10 as indicated.

Please add new claims 11 and 12.

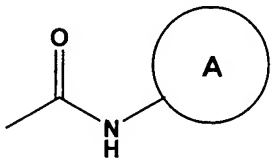
Claim 1 (currently amended): A compound of formula **(I)** or a salt, solvate or pro-drug thereof:



(I)

wherein:

one of $\mathbf{R^1}$ and $\mathbf{R^2}$ is a group **(IA)**:



(IA)

and the other of $\mathbf{R^1}$ and $\mathbf{R^2}$ is C_{1-4}alkoxy~~; wherein this R^1 or R^2 is~~ optionally substituted on carbon by one or more groups selected from R^5 ; ~~and wherein if said heterocyclyl contains an –NH– moiety that nitrogen is optionally substituted by C_{1-4}alkyl;~~

Ring A is pyridin-2-yl ~~pyridin-2yl~~ or thiazol-2-yl; wherein said pyridin-2-yl or thiazol-2-yl is optionally substituted on carbon by one or more groups selected from R^6;

one of $\mathbf{R^3}$ and $\mathbf{R^4}$ is hydrogen and the other is selected from hydrogen, C_{1-4}alkyl, C_{1-4}alkoxy, carbocyclyl, heterocyclyl, carbocyclyloxy and heterocyclyloxy; wherein R^3 and R^4 are independently optionally substituted on carbon by one or more groups selected from R^7; and wherein if R^3 or R^4 is heterocyclyl, said heterocyclyl contains an -NH- moiety, and the ~~that~~ nitrogen of the -NH-moiety is optionally substituted by C_{1-4}alkyl;

$\mathbf{R^6}$ is selected from halo, carboxy and C_{1-4}alkyl;

$\mathbf{R^5}$ and $\mathbf{R^7}$ ~~is~~ are independently selected from halo, C_{1-4}alkyl, C_{1-4}alkoxy, *N*-(C_{1-4}alkyl)amino, *N,N*-(C_{1-4}alkyl)$_2$amino, carbocyclyl, heterocyclyl, carbocyclyloxy, heterocyclyloxy and carbocyclylidenyl; wherein $\mathbf{R^5}$ and $\mathbf{R^7}$ are independently optionally substituted on carbon by one or more groups selected from $\mathbf{R^8}$; and wherein if R^5 and/or R^7 is heterocyclyl, said heterocyclyl contains an -NH- moiety, and the ~~that~~ nitrogen of the -NH-moiety is optionally substituted by C_{1-4}alkyl; and

$\mathbf{R^8}$ is selected from halo, carboxy, methyl, ethyl, methoxy, ethoxy, methylamino, ethylamino, dimethylamino, diethylamino and *N*-methyl-*N*-ethylamino~~;~~

~~or a salt, solvate or pro-drug thereof~~.

Claim 2 (cancelled).

Claim 3 (currently amended): A compound according to Claim ~~2~~ 1 or a salt, solvate or pro-drug thereof, wherein Ring **A** in the group **(IA)** is substituted by carboxy and the C_{1-4}alkoxy group is substituted on carbon by one or more groups selected from $\mathbf{R^5}$.

Claim 4 (currently amended): A compound according to Claim 3 or a salt, solvate or pro-drug thereof, wherein $\mathbf{R^5}$ is selected from carbocyclyl optionally substituted by one or more groups selected from $\mathbf{R^8}$.

Claim 5 (currently amended): A compound according to Claim 1 or a salt, solvate or pro-drug thereof,wherein one of $\mathbf{R^3}$ and $\mathbf{R^4}$ is hydrogen and the other is C_{1-4}alkyl.

Claim 6 (currently amended): A compound according to Claim 1 or a salt, solvate or pro-drug thereof, selected from:

2-(2-Chlorobenzyloxy)-4-[*N*-5-carboxythiazol-2-yl)carbamoyl]-6-methylquinoline;

2-(2-Chlorobenzyloxy)-4-[*N*-5-carboxythiazol-2-yl)carbamoyl]-quinoline;

2-(2-Chlorobenzyloxy)-4-[*N*-5-carboxypyrid-2-yl)carbamoyl]-6-methylquinoline;

2-(2-Chlorobenzyloxy)-4-[*N*-5-carboxypyrid-2-yl)carbamoyl]-quinoline;

2-[*N*-5-carboxyprid-2-yl)carbamoyl]-4-(2-methylbenzyloxy)-quinoline; and

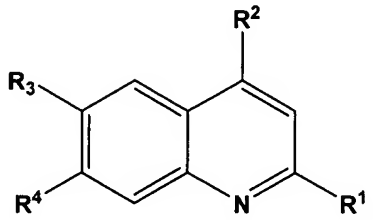
2-(1-methylpropoxy)-4-[*N*-(5-carboxythiazol-2-yl)carbamoyl]-quinoline~~;~~

~~or a salt, solvate or pro-drug thereof~~.

Claim 7 (previously presented): A pharmaceutical composition comprising a compound according to any one of Claims 1 and 3 to 6, or a salt, pro-drug or solvate thereof, together with a pharmaceutically acceptable diluent or carrier.

Claim 8 (previously presented): A method of treating diabetes and/or obesity by administering an effective amount of a compound according to Claim 1 or a salt, solvate or pro-drug thereof, to a mammal in need of such treatment.

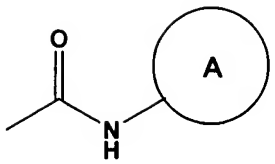
Claim 9 (currently amended): A process for preparing a compound of formula **(I)** or a salt, solvate or pro-drug thereof,:



(I)

wherein:

one of $\mathbf{R^1}$ and $\mathbf{R^2}$ is a group **(IA)**:



(IA)

and the other of $\mathbf{R^1}$ and $\mathbf{R^2}$ is C_{1-4}alkoxy~~; wherein this R^1 or R^2 is~~ optionally substituted on carbon by one or more groups selected from R^5; ~~and wherein if said heterocyclyl contains an -NH- moiety that nitrogen is optionally substituted by C_{1-4}alkyl;~~

Ring A is pyridin-2-yl ~~pyridin-2yl~~ or thiazol-2-yl; wherein said pyridin-2-yl or thiazol-2-yl is optionally substituted on carbon by one or more groups selected from R^6;

one of $\mathbf{R^3}$ and $\mathbf{R^4}$ is hydrogen and the other is selected from hydrogen, C_{1-4}alkyl, C_{1-4}alkoxy, carbocyclyl, heterocyclyl, carbocyclyloxy and heterocyclyloxy; wherein R^3 and R^4 are independently optionally substituted on carbon by one or more groups selected from R^7; and wherein if R^3 or R^4 is heterocyclyl, said heterocyclyl contains an -NH- moiety, and the ~~that~~ nitrogen of the -NH- moiety is optionally substituted by C_{1-4}alkyl;

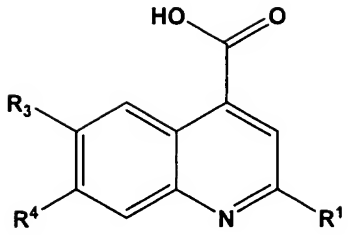
$\mathbf{R^6}$ is selected from halo, carboxy and C_{1-4}alkyl;

$\mathbf{R^5}$ and $\mathbf{R^7}$ ~~is~~ are independently selected from halo, C_{1-4}alkyl, C_{1-4}alkoxy, *N*-(C_{1-4}alkyl)amino, *N,N*-(C_{1-4}alkyl)$_2$amino, carbocyclyl, heterocyclyl, carbocyclyloxy, heterocyclyloxy and carbocyclylidenyl; wherein $\mathbf{R^5}$ and $\mathbf{R^7}$ are independently optionally substituted on carbon by one or more $\mathbf{R^8}$; and wherein if R^5 and/or R^7 is heterocyclyl, said heterocyclyl contains an -NH- moiety, and the ~~that~~ nitrogen of the -NH-moiety is optionally substituted by C_{1-4}alkyl; and

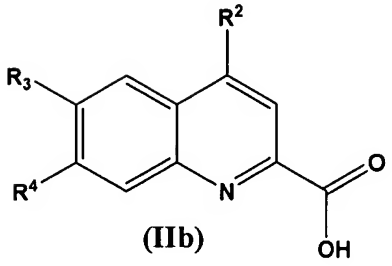
$\mathbf{R^8}$ is selected from halo, carboxy, methyl, ethyl, methoxy, ethoxy, methylamino, ethylamino, dimethylamino, diethylamino and *N*-methyl-*N*-ethylamino~~;~~

~~or a salt, solvate or pro-drug thereof,~~ which process comprises:

Process 1): reacting an acid of formula **(IIa)** or **(IIb)**:

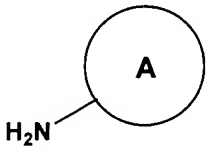


(IIa)



(IIb)

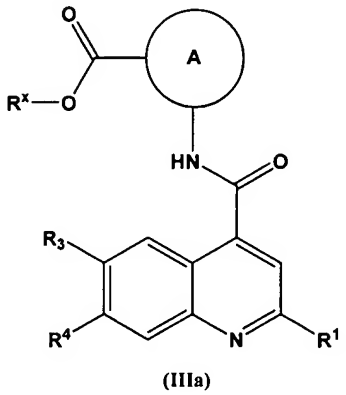
or an activated derivative thereof with a compound of formula **(III)**



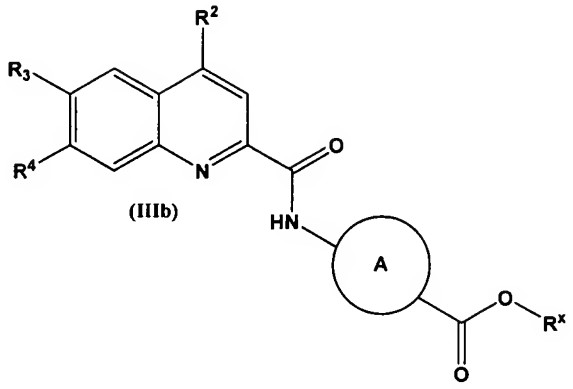
(III)

or

Process 2): for compounds of formula **(I)** wherein R^6 is carboxy, deprotecting a compound of formula **(IIIa)** or **(IIIb)**:



(IIIa)



(IIIb)

wherein ~~R^xC(O)O-~~ R^xOC(O)- is an ester group and R^x is selected from C_{1-6} alkyl and benzyl;

and thereafter ~~if necessary or desirable:~~ optionally,

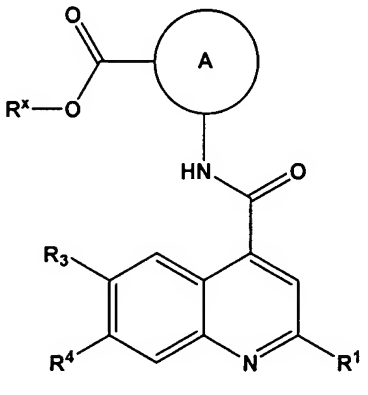
i) converting a compound of the formula (I) into another compound of the formula **(I)**;

and/or

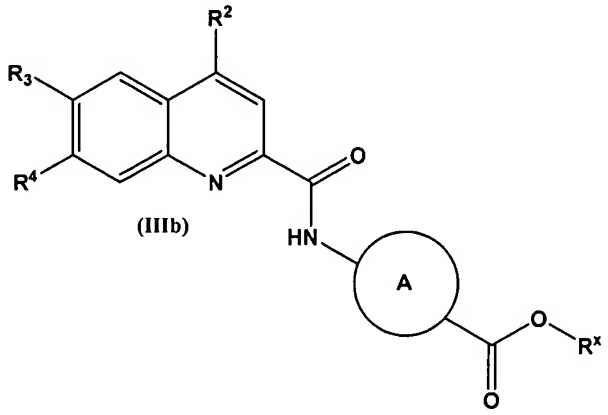
ii) removing any protecting groups; and/or

iii) forming a salt, solvate or pro-drug thereof.

Claim 10 (currently amended): A compound of formula (IIIa) or a compound of formula (IIIb):



(IIIa)



(IIIb)

wherein ~~R^xC(O)O-~~ R^xOC(O)- is an ester group and R^x is selected from C_{1-6} alkyl and benzyl;

R^1 and **R^2** are C_{1-4}alkoxy, ~~is C_{1-4}alkoxy; wherein this R^1 or R^2 is~~ optionally substituted on carbon by one or more groups selected from R^5; ~~and wherein if said heterocyclyl contains an –NH– moiety that nitrogen is optionally substituted by C_{1-4}alkyl;~~

Ring A is pyridin-2-yl ~~pyridin-2yl~~ or thiazol-2-yl; wherein said pyridin-2-yl or thiazol-2-yl is optionally substituted on carbon by one or more groups selected from R^6;

one of $\mathbf{R^3}$ and $\mathbf{R^4}$ is hydrogen and the other is selected from hydrogen, C_{1-4}alkyl, C_{1-4}alkoxy, carbocyclyl, heterocyclyl, carbocyclyloxy and heterocyclyloxy; wherein R^3 and R^4 are independently optionally substituted on carbon by one or more groups selected from R^7; and wherein if R^3 or R^4 is heterocyclyl, said heterocyclyl contains an -NH- moiety, and the ~~that~~ nitrogen of the -NH- moiety is optionally substituted by C_{1-4}alkyl;

$\mathbf{R^6}$ is selected from halo, carboxy and C_{1-4}alkyl;

$\mathbf{R^5}$ and $\mathbf{R^7}$ ~~is~~ are independently selected from halo, C_{1-4}alkyl, C_{1-4}alkoxy, *N*-(C_{1-4}alkyl)amino, *N,N*-(C_{1-4}alkyl)$_2$amino, carbocyclyl, heterocyclyl, carbocyclyloxy, heterocyclyloxy and carbocyclylidenyl; wherein $\mathbf{R^5}$ and $\mathbf{R^7}$ are independently optionally substituted on carbon by one or more $\mathbf{R^8}$; and wherein if R^5 and/or R^7 is heterocyclyl, said heterocyclyl contains an -NH- moiety, and the ~~that~~ nitrogen of the -NH- moiety is optionally substituted by C_{1-4}alkyl; and

$\mathbf{R^8}$ is selected from halo, carboxy, methyl, ethyl, methoxy, ethoxy, methylamino, ethylamino, dimethylamino, diethylamino and *N*-methyl-*N*-ethylamino.

Claim 11 (new): The process of claim 9, wherein R^x is selected from methyl and ethyl.

Claim 12 (new): The compound of claim 10, wherein R^x is selected from methyl and ethyl.